



Europäisches Patentamt
European Patent Office
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⑪ Publication number:

0 218 736
A1

⑫

EUROPEAN PATENT APPLICATION

⑬ Application number: 85112797.7

⑬ Int. Cl. C07D 409/06, C07D 409/14,
A01N 43/50

⑭ Date of filing: 09.10.85

⑮ Date of publication of application:
22.04.87 Bulletin 87/17

⑯ Designated Contracting States:
AT BE CH DE FR GB IT LI LU NL SE

⑭ Applicant: NIHON NOHYAKU CO., LTD.
1-2-6, Nihonbashi
Chuo-ku Tokyo(JP)

⑯ Inventor: Soe, Akira
6-2-312, Salwalicho
Yamatotakada-shi(JP)
Inventor: Kanno, Hideo
2-2-708, Shirakawa-3-chome
Ibaraki-shi(JP)
Inventor: Hasegawa, Nobu
1-10, Uedanishimachi
Nishinomya-shi(JP)
Inventor: Miyagi, Yukio
7-23-816, Nankonaka-4-chome
Suminoo-ku Osaka(JP)
Inventor: Nishimura, Akira
5-6, Hondacho
Kawachinagano-shi(JP)
Inventor: Konaka, Shigeo
9-4-213, Takewashi-4-chome
Habikino-shi(JP)
Inventor: Ohmi, Tetsuto
1-28, Nishinoyamacho
Kawachinagano-shi(JP)
Inventor: Munechika, Yukimi
3-8-1, Toge
Hashimoto-shi(JP)
Inventor: Uchida, Matazaemon
14-22, Daishicho
Kawachinagano-shi(JP)
Inventor: Ikeda, Kenichi
13-13, Tsuganodal-1-chome
Chiba-shi(JP)

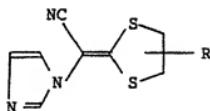
⑯ Representative: Patentanwälte Grünnecker,
Kinkeldey, Stockmair & Partner
Maximilianstrasse 88
D-8000 München 22(DE)

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⑯ A novel ketene S,S-acetal derivative, a process for manufacturing thereof and a method for curing mycosis by administering it.

⑤ A novel ketene S,S-acetal derivative which is useful as an antimycotic agent and an agricultural chemical of fungicidal, plant growth regulating or

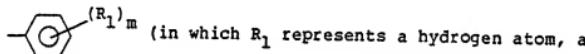
insecticidal properties represented by the general formula (I):



(I)

(in which R, represents a hydrogen atom, a wherein R represents a hydrogen atom; an alkyl group having 1 to 8 carbon atoms; a cycloalkyl group having 3 to 6 carbon atoms; a methylene group; a lower alkenyl group; a lower alkyl group

substituted by a halogen atom, a cyano group, a lower alkoxy group, a lower alkylthio group; a carbamoyl group, an acyl group, or an alkenyloxy group; a phenyl group represented by



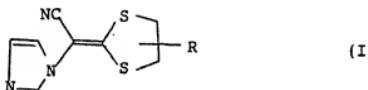
halogen atom, a straight or branched chain lower alkyl group, a lower alkoxy group which may be substituted by one or more halogen atoms, a phenoxy group or a methylenedioxy group, and m represents an integer of 1 to 3); a benzyl group; a methylenedioxybenzyl group; a phenoxyalkyl group; a phenoxyalkyl group substituted by a halogen atom; a naphthyl group; or a substituted or unsubstituted pyridyl group.

A NOVEL KETENE FOR S,S-ACETAL DERIVATIVE, A PROCESS FOR MANUFACTURING THEREOF AND A METHOD FOR CURING MYCOSIS BY ADMINISTERING IT

BACKGROUND OF THE INVENTION

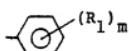
This invention relates to a novel ketene S,S-acetal derivatives which are useful as antimycotic agents and agricultural chemicals.

More particularly, this invention relates to ketene S,S-acetal derivatives represented by the general formula (I):



wherein R represents a hydrogen atom; an alkyl group having 1 to 8 carbon atoms; a cycloalkyl group having 3 to 6 carbon atoms; a methylene group; a lower alkenyl group; a lower alkyl group

15 substituted by a halogen atom, a cyano group, a lower alkoxy group, a lower alkylthio group, a carbamoyl group, an acyl group, or an alkenyloxy group; a phenyl group represented by



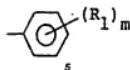
(in which R₁ represents a hydrogen atom, a halogen atom, a straight or branched chain lower alkyl group, a lower alkoxy group which may be substituted by one or more halogen atoms, a phenoxy group or a methylenedioxy group, and m represents an integer of 1 to 3); a benzyl group; a methylenedioxybenzyl group; a phenoxyalkyl group; a phenoxyalkyl group substituted by a halogen atom; a naphthyl group; or a substituted or unsubstituted pyridyl group.

cides, whereby this invention has been accomplished.

The substituent R in the above general formula (I) includes hydrogen atom; straight or branched chain alkyl groups such as methyl, ethyl, n-propyl, i-propyl, n-butyl, i-butyl, s-butyl, t-butyl, n-pentyl, i-pentyl, neo-pentyl, n-hexyl, n-octyl and the like; cycloalkyl groups such as cyclopropyl, cyclohexyl and the like; methylene group; lower alkenyl groups such as vinyl, allyl and the like; lower alkyl groups substituted by a halogen atom such as chloromethyl, dichloroethyl and the like; lower alkyl groups substituted by a cyano group such as cyanomethyl and the like; lower alkyl groups substituted by a lower alkoxy such as methoxymethyl, methoxyethyl and the like or a lower alkylthio group such as methythiomethyl and the like; lower alkyl group substituted by carbamoyl group such as carbamoylmethyl and the like; lower alkyl groups substituted by an acyl group; lower alkyl groups substituted by an alkenyloxy groups such as acryloylmethyl and the like; phenyl groups represented by

SUMMARY OF INVENTION

The present inventors have devoted themselves to research in order to invent a novel ketene S,S-acetal derivative, and have consequently found that compounds represented by the general formula (I) are novel compounds not described in literatures and are useful as antimycotic agents and agricultural chemicals; in particular, not only fungicides and plant growth regulators but also insecti-



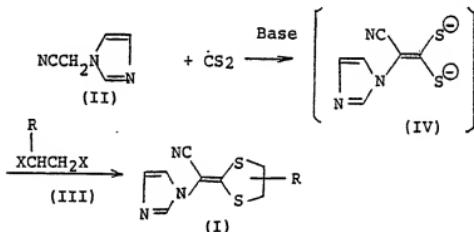
(in which R_1 represents a hydrogen atom, a halogen atom, a straight or branched chain lower alkyl group, a lower alkoxy group which may be substituted by one or more halogen atoms, a phenoxy group or a methylenedioxy group, and m represents an integer of 1 to 3); a benzyl group; a methylenedioxybenzyl group; a phenoxyalkyl group such as phenoxyethyl and the like; a phenoxyalkyl group substituted by a halogen atom; naphthyl

group; substituted or unsubstituted pyridyl group; etc.

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DETAILED DESCRIPTION OF INVENTION

The compound represented by the general formula (I) of this invention can be synthesized, for example, by the process shown below:



wherein R is as defined above, and X represents a halogen atom, a mesyloxy or a tosylxy group. That is to say, the compound represented by the general formula (I) can be obtained by reacting 1-cyanomethylimidazole represented by the structural formula (II) with carbon disulfide in the presence of a base and a solvent to form an intermediate represented by the structural formula (IV), and reacting the intermediate with a compound represented by the general formula (III) without isolating the same.

As the solvent usable in the invention, any solvent may be used so long as it does not inhibit the progress of the reaction, and there can be exemplified, for example, alcohols such as methanol, ethanol, isopropanol and the like, dimethylsulfoxide, dimethylformamide, hexamethylenephosphoramide, water, etc. These solvents can be used alone or as a mixture thereof.

As the base usable in this invention, there can be exemplified sodium carbonate, potassium carbonate, sodium hydrogencarbonate, potassium

hydrogencarbonate, sodium hydroxide, potassium hydroxide, potassium t-butoxide, etc. These can be used in solid state or in solution.

35 Although it is sufficient that the reaction temperature is selected in the range of 0° to 100°C, it is particularly preferable to carry out the reaction at a temperature near room temperature.

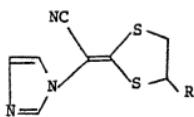
It is sufficient that the reaction time is properly selected in the range of 0.5 to 24 hours.

40 It is sufficient that the amount of the base used is selected in the range of 2 to 4 moles per mole of 1-cyanomethylimidazole represented by the structural formula (IV).

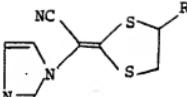
45 It is sufficient that after completion of the reaction, the reaction solution is treated in the usual way. For example, the reaction product is extracted and separated with a suitable solvent and then can be purified by recrystallization or column chromatography.

50 The compound represented by the general formula (I) is often obtained as a mixture of two kinds of the geometrical isomers shown below.

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(Z isomer)



(E isomer)

The aforesaid mixture of the Z and E isomers can often be isolated into the two isomers by a suitable separation method, for example, recrystallization, chromatography or the like.

10 This invention includes the geometrical isomers, i.e., the E and Z isomers, and all mixtures of the two isomers in any ratio.

15 Typical examples of the compounds represented by the general formula (I) are shown in Table 1, but this invention is not limited thereto.

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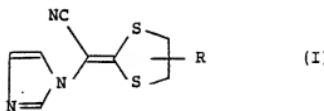
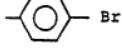
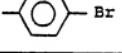
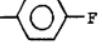


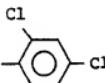
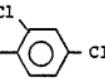
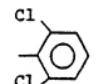
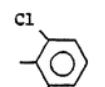
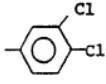
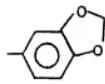
Table 1

Com- ound No.	R	Physical property, melting point, refractive index, NMR value (TMS/CDCl ₃) -
1	H	Melting point 126.5°C
2	CH ₃	Melting point 97.6°C
3	C ₂ H ₅	n_D^{17} 1.6246
4	n-C ₃ H ₇	n_D^{27} 1.6065
5	i-C ₃ H ₇	Melting point 86.7°C (Z isomer)
6	i-C ₃ H ₇	Melting point 55.8°C (E isomer)
7	n-C ₄ H ₉	n_D^{27} 1.5830
8	i-C ₄ H ₉	Melting point 73.3°C (Z isomer)
9	i-C ₄ H ₉	Melting point 118.1°C (F isomer)

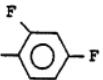
Com- ound No.	R	Physical property, melting point, refractive index, NMR value (TMS/CDCl ₃)
10	s-C ₄ H ₉	n _D ¹⁶ 1.6011 (Z isomer)
11	s-C ₄ H ₉	n _D ¹⁶ 1.6089 (E isomer)
12	t-C ₄ H ₉	Melting point 151.7°C
13	n-C ₅ H ₁₁	n _D ^{16.5} 1.5931 (Z isomer)
14	n-C ₅ H ₁₁	n _D ^{16.5} 1.5949 (E isomer)
15	i-C ₅ H ₁₁	Melting point 74.3°C (Z isomer)
16	i-C ₅ H ₁₁	Melting point 111.4°C (E isomer)
17	neo-C ₅ H ₁₁	Melting point 96.2°C (Z isomer)
18	neo-C ₅ H ₁₁	Melting point 107.7°C (E isomer)
19	n-C ₆ H ₁₃	n _D ¹⁷ 1.5908 (Z isomer)
20	n-C ₆ H ₁₃	Melting point 48.2°C (E isomer)
21		Melting point 113.9°C (Z isomer)
22		Melting point 103.8°C (E isomer)
23	=CH ₂	Melting point 115.7°C
24	-CH=CH ₂	n _D ²⁷ 1.6422

Com- ound No.	R	Physical property, melting point, refractive index, NMR value (TMS/CDCl ₃)
25	-CH ₂ Cl	Melting point 96.2°C
26	-CHClCH ₂ Cl	n _D ¹⁶ 1.6103
27	-CH ₂ CN	n _D ¹⁸ 1.6167
28	-CH ₂ CH ₂ OCH ₂ CH ₃	n _D ²⁴ 1.6009
29	-CH ₂ SCH ₃	n _D ^{13.5} 1.6412
30	-CH ₂ CONH ₂	Viscous oily substance 3.64 (m, 2H), 4.20 (m, 3H), 5.27 (br, 2H) 6.96, 7.05, 7.52 (1H on hetero ring for each)
31	-CH ₂ CH ₂ COCH ₃	n _D ¹⁸ 1.6086
32	-CH ₂ O ₂ CCH=CH ₂	n _D ¹⁸ 1.6082
33		n _D ²⁴ 1.6417
34		Melting point 119.4°C (Z isomer)
35		Melting point 141.5°C (E isomer)
36		n _D ²⁴ 1.6083

No.	R	Physical property, melting point, refractive index, NMR value (TMS/CDCl ₃)
37		Viscous oily substance (Z isomer) 3.54-4.20 (m, 2H), 5.69 (dd, 1H) 7.00-7.75 (m, 7H)
38		Viscous oily substance (E isomer) 3.45-4.10 (m, 2H), 5.76 (dd, 1H) 7.00-7.85 (m, 7H)
39		Viscous oily substance (z isomer) 3.6-4.0 (m, 2H), 5.22 (dd, 1H) 6.9-7.81 (m, 7H)
40		Melting point 148.8°C (E isomer)
41		Viscous oily substance (Z isomer)
42		Melting point 119°C (E isomer)
43		Viscous Oily substance 3.6-3.9 (m, 2H), 5.1-5.5 (m, 1H) 6.9-7.8 (m, 7H)
44		n_D^{18} 1.6313 (Z isomer)

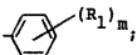
Com- ound No.	R	Physical property, melting point, refractive index, NMR value (TMS/CDCl ₃)
52		Melting point 110.5°C (Z isomer)
53		Melting point 100.4°C (E isomer)
54		Viscous oily substance (Z isomer)
55		Viscous oily substance (E isomer) 3.42 (dd, 1H), 4.43 (t, 1H), 6.16 (dd, 1H), 7.18-7.40 (m, 3H), 6.98, 7.08, 7.56 (1H for each, H on azole ring)
56		Viscous oily substance 3.6-3.9 (m, 2H), 5.0-5.4 (m, 1H), 6.9-7.8 (m, 6H)
57		Viscous oily substance 3.64, 3.72 (d, 2H for each), 5.11, 5.19 (t, 1H for each), 5.92, 5.95 (s, 2H for each), 6.60-6.95 (m, 3H), 7.00, 7.08, 7.53 (1H for each, H on azole ring)

Compound No.	R	Physical property, melting point, refractive index, NMR value (TMS/CDCl ₃)
58		Melting point 187.5°C (Z isomer)
59		Melting point 214.0°C (E isomer)
60	-CH ₂ -	Melting point 102.7°C
61	-CH ₂ -	n _D ¹⁷ 1.6258
62	-CH ₂ O-	n _D ¹⁸ 1.6315
63	-CH ₂ O-	n _D ¹⁸ 1.6213
64		Melting point 135.0°C
65		Viscous oily substance 3.65-4.38 (m, 2H), 5.22-5.60 (m, 1H), 7.00-7.95 (m, 6H), 8.54-8.76 (m, 1H)

Com- ound No.	R	Physical property, melting point, refractive index, NMR value (TMS/CDCl ₃)
66		Viscous oily substance 3.75 (m, 2H), 5.20 (m, 1H), 6.80-7.80 (m, 7H)
67		$n_D^{18.5}$ 1.6063
68		Viscous oily substance (Z isomer) 3.83 (m, 2H), 5.46 (dd, 1H), 6.70-7.80 (m, 6H)
69		Viscous oily substance (E isomer) 3.75 (m, 2H), 5.52 (dd, 1H), 6.60-7.80 (m, 6H)
70		$n_D^{17.5}$ 1.6358 (E isomer)
71		$n_D^{17.5}$ 1.6458 (Z isomer)
72		Viscous oily substance (Z isomer) 3.83 (m, 2H), 5.59 (dd, 1H), 6.75-7.70 (m, 6H)

Compound No.	R	Physical property, melting point, refractive index, NMR value (TMS/CDCl ₃)
73		m.p. 146-152°C (E isomer)
74		Viscous oily substance (Z isomer) 3.38 (m, 2H), 5.42 (dd, 1H), 6.90-7.70 (m, 6H)
75		m.p. 114-117°C (E isomer)
76		$n_{D}^{22.0}$ 1.6401
77		$n_{D}^{22.0}$ 1.6262
78		$n_{D}^{25.5}$ 1.6432
79		$n_{D}^{25.5}$ 1.6332

Preferred compounds are these in which R ⁵⁰ represents a lower alkyl group, a lower alkyl group substituted by a lower alkoxy or alkylthio group, or



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in which R₁ represents a halogen atom, especially chlorine or bromine atom, or a straight or branched chain lower alkyl group, especially methyl, and m represents an integer of 1 to 3.

Examples of this invention are shown below.

Example 1

Synthesis of 2-(1-imidazolyl)-2-(4-isobutyl-1,3-dithiolan-2-ylidene)acetonitrile (Compound Nos. 8 and 9)

To a mixed solution of 0.55 g (0.005 mole) of 1-cyanomethylimidazole, 0.4 g (0.005 mole) of carbon disulfide and 10 ml of dimethyl sulfoxide was added 0.8 g (0.014 mole) of potassium hydroxide powder with stirring, and the reaction was carried out at room temperature for 1 hour. Then, 1.5 g (0.006 mole) of 1,2-dibromo-4-methylpentane was added dropwise with stirring, and the resulting solution was subjected to reaction for 2 hours. After completion of the reaction, 20 ml of water was added to the reaction solution, after which the resulting mixture was subjected to extraction with ethyl acetate, and the organic layer was washed with water and dried. The solvent was distilled off and the residue was purified by silica gel chromatography to obtain 0.45 g of the Z isomer and 0.3 g of the E isomer individually in the form of colorless crystals.

The Z isomer (Compound No. 8): melting point 73.3°C, yield 34%.

The E isomer (Compound No. 9): melting point 118.1°C, yield 23%.

Example 2

Synthesis of 2-(1-imidazolyl)-2-(4-chloromethyl-1,3-dithiolan-2-ylidene)acetonitrile (Compound No. 25).

To a mixed solution of 0.55 g (0.005 mole) of 1-cyanomethylimidazole, 0.4 g (0.005 mole) of carbon disulfide and 10 ml of dimethyl sulfoxide was added 0.8 g (0.014 mole) of potassium hydroxide powder with stirring, and the reaction was carried out at room temperature for 1 hour. Then, 1.4 g (0.006 mole) of 1,2-dibromo-3-chloropropane was added dropwise with stirring, and the resulting solution was subjected to reaction for 2 hours. After completion of the reaction, 20 ml of water was

added to the reaction solution, after which the resulting mixture was subjected to extraction with ethyl acetate, and the organic layer was washed with water and dried. The solvent was distilled off and the residue was purified by silica gel chromatography and recrystallized from ethyl acetate-n-hexane to obtain 0.8 g of the desired compound in the form of colorless crystals: melting point 96.2°C, yield 62%.

Example 3

Synthesis of 2-(1-imidazolyl)-2-(4-methylidene-1,3-dithiolan-2-ylidene)acetonitrile (Compound No. 23)

In 10 ml of tetrahydrofuran were dissolved 0.52 g (0.002 mole) of the 2-(1-imidazolyl)-2-(4-chloromethyl-1,3-dithiolan-2-ylidene)acetonitrile obtained in Example 2 and 0.31 g of 1,8-diazabicyclo[5.4.0]7-undecene, and the reaction was carried out with heating under reflux for 1 hour. After the reaction solution was allowed to cool, the deposited salt was separated by filtration and the filtrate was concentrated to obtain crude crystals, which were then recrystallized from ethyl acetate-n-hexane to obtain 0.35 g of the desired compound in the form of crystals: melting point 115.7°C, yield 79%.

Example 4

Synthesis of 2-(1-imidazolyl)-2-[4-(2,4-dichlorophenyl)-1,3-dithiolan-2-ylidene]acetonitrile (Compound Nos. 52 and 53)

To a mixed solution of 0.55 g (0.005 mole) of 1-cyanomethylimidazole, 0.4 g (0.005 mole) of carbon disulfide and 10 ml of dimethyl sulfoxide was added 0.8 g (0.014 mole) of potassium hydroxide powder with stirring, and the reaction was carried out at room temperature for 1 hour. Then, 2.0 g (0.006 mole) of 2',4'-dichloro-1,2-dibromoethoxybenzene was added dropwise with stirring, and the resulting solution was subjected to reaction for 2 hours. After completion of the reaction, 20 ml of water was added to the reaction solution, and the resulting mixture was subjected to extraction with ethyl acetate, and the organic layer was washed with water and dried. The solvent was distilled off and the residue was purified by silica gel chromatography to obtain 0.25 g of the Z isomer

and 0.5 g of the E isomer individually in the form of a yellow viscous substance: yield (total yield of the Z and E isomers) 42%, melting points 110.5°C (the Z isomer) and 100.4°C (the E isomer).

Example 5

Synthesis of 2-(1-imidazolyl)-2-[4-(2-isopropyl-phenyl)-1,3-dithiolan-2-ylidene]acetonitrile (Compound No. 47)

To mixed solution of 0.55 g (0.005 mole) of 1-cyanoethylimidazole, 0.4 (0.005 mole) of carbon disulfide and 10 ml of dimethylformamide was added 0.8 g (0.014 mole) of potassium hydroxide powder, and the reaction was carried out with stirring at room temperature for 1 hour. Then, 1.8 g (0.006 mole) of 2'-isopropyl-1,2-dibromoethylbenzene was added dropwise with stirring, and the resulting solution was subjected to reaction for another 2 hours. After completion of the reaction, 20 ml of water was added to the reaction solution, and the resulting mixture was subjected to extraction with ethyl acetate, and the organic layer was washed with water and dried. The solvent was distilled off and the residue was purified by silica gel chromatography to obtain the desired compound in the form of a light-yellow oily substance: $n_D^{25} 1.8196$, yield 34%.

Example 6

Synthesis of 2-(1-imidazolyl)-2-[4-(2-chlorophenyl)-1,3-dithiolan-2-ylidene]acetonitrile (Compound Nos. 34 and 35)

To a mixed solution of 2.2 g (0.02 mole) of 1-cyanoethylimidazole, 1.60 g (0.02 mole) of carbon disulfide and 10 ml of dimethyl sulfoxide was added 3.0 g (0.05 mole) of potassium hydroxide powder, and the reaction was carried out with stirring at room temperature for 1 hour. Then, 4.2 g (0.02 mole) of 2'-chloro-(1,2-dichloroethyl)benzene was added dropwise with stirring, and the resulting solution was subjected to reaction for another 2 hours. After completion of the reaction, 20 ml of water was added to the reaction solution, after which the resulting mixture was subjected to extraction with ethyl acetate, and the organic layer was washed with water and dried. The solvent was distilled off and the residue was purified by silica gel chromatography to obtain 2.0 g of the Z isomer and 2.4 g of the E isomer individually in the form of light-yellow crystals.

The Z isomer: melting point 119.4°C, yield 31%.

The E isomer: melting point 141.5°C, yield 38%.

Example 7

Synthesis of 2-(1-imidazolyl)-2-[4-(2-chlorophenyl)-1,3-dithiolan-2-ylidene]acetonitrile (Compound Nos. 34 and 35)

To a mixed solution of 2.2 g (0.02 mole) of 1-cyanoethylimidazole, 1.60 g (0.02 mole) of carbon disulfide and 10 ml of dimethyl sulfoxide was added 3.0 g (0.05 mole) of potassium hydroxide powder, and the reaction was carried out with stirring at room temperature for 1 hour. Then, 6.0 g (0.02 mole) of 2'-chloro-(1,2-dibromoethyl)benzene was added dropwise with stirring, and the resulting solution was subjected to reaction for another 2 hours. After completion of the reaction, 20 ml of water was added to the reaction solution, after which the resulting mixture was subjected to extraction with ethyl acetate, and the organic layer was washed with water and dried. The solvent was distilled off and the residue was purified by silica gel chromatography to obtain 0.3 g of the Z isomer and 1.3 g of the E isomer individually in the form of light-yellow crystals.

The Z isomer: melting point 119.4°C, yield 5%.
The E isomer: melting point 141.5°C, yield 20%.

Example 8

Synthesis of 2-(1-imidazolyl)-2-[4-(2-chlorophenyl)-1,3-dithiolan-2-ylidene]acetonitrile (Compound Nos. 34 and 35)

To a mixed solution of 2.2 g (0.02 mole) of 1-cyanoethylimidazole, 1.60 g (0.02 mole) of carbon disulfide and 10 ml of dimethyl sulfoxide was added 3.0 g (0.05 mole) of potassium hydroxide powder, and the reaction was carried out with stirring at room temperature for 1 hour. Then, 6.6 g (0.02 mole) of 2'-chloro-(1,2-dimesyloxyethyl)benzene was added dropwise with stirring, and the resulting solution was subjected to reaction for another 2 hours. After completion of the reaction, 20 ml of water was added to the reaction solution, after which the resulting mixture was subjected to extraction with ethyl acetate, and the organic layer was washed with water and dried. The solvent was distilled off and the residue was purified by silica gel chromatography to obtain 1.3 g of the Z isomer and 1.8 g of the E isomer individually in the form of light-yellow crystals.

The Z isomer: melting point 119.4°C, yield 20%.

The E isomer: melting point 141.5°C, yield 28%.

The compounds of this invention are antimycotic agents useful for preventing fungal infection of human beings and animals. For example, these compounds can be used for curing mycoses such as local mycotic infection, mucosal mycotic infection, systemic mycotic infection and the like which are caused by Dermatophytes such as *Microsporum*, *Epidermophyton*, *Trichophyton* and the like and *Candida*.

The compounds of this invention can be mixed with conventional chemotherapeutically acceptable diluents or carriers and if desired, other excipients, and can be used in pharmaceutical forms such as solutions, creams, suppositories, ointments, tablets, etc.

When used as antimycotic agents, the compounds of this invention can be used as local liniments in pharmaceutical forms such as creams, ointments, solution, etc. When they are used in the form of an endermic solution, their practical concentration is considered to be suitably 0.1% or more.

The present drugs may be used, of course, in an admixture with other antibacterial agents such as amphotericin B, nystatin, trichomycin, variolin, clotrimazole and the like.

Further, the compounds of this invention are useful as agricultural and horticultural fungicides. For example, they are very effective against various phytopathogenic diseases, e.g., rice blast - (*Phricularia oryzae*); powdery mildew of barley and wheat (*Erysiphe graminis*), and other powdery mildews of various host plants such as that of cucumber (*Sphaerotheca fuliginea*), that of apple - (*Podosphaera leucotricha*) and that of grape - (*Uncinia necator*); rust of wheat (*Puccinia recondita*); Crown rust of oats (*Puccinia coronata*) and rust of other host plants; late blight of tomato - (*Phytophthora capsici*) and phytophthora rot of other host plants; etc.

When the compound of this invention is used as active ingredient in an agricultural and a horticultural fungicide, the fungicide is prepared into a formulation suitable for use in a conventional manner as an agricultural chemical. For example, the fungicide is prepared in the form of dust, granules, fine granules, wettable powder, emulsifiable concentrate, oily solution, aerosol, floating dust, fumigants, preparations suitable for vaporizing the active ingredient by heat or other physical means, tablets or the like by mixing the compound of the invention with adjuvants, and is applied to stalks and leaves of vegetables, flowering plants, crops for industrial use, fruit trees, other trees and the

like as it is or after diluted to a suitable volume with water. Although in this invention the dosage of the active ingredient varies depending on the kind of the compound, plant to be treated, the way of using, and the like, it can be selected in the range of 5 to 500 g per 10 acres.

When applied, the compound of this invention can also be used in admixture with or in combination with other agricultural chemicals, fertilizers, plant nutrients and the like which can be used similarly to the compound.

For example, when a phytopathogenic disease is controlled by using an agricultural and horticultural fungicide comprising the compound of this invention as active ingredient, the fungicide can be made into a multiple-purpose preventing and curing agent by mixing therewith an agent for preventing and curing other diseases and/or vermin insects which break out simultaneously with aforesaid disease.

Next, some Test Examples and Formulation Examples are given below in order to demonstrate the usefulness of the compounds of this invention, but this invention is not intended to be limited thereto.

Test Example 1

30 Test for antifungal activity against *Trichophyton mentagrophytes*

In 1 liter of water were dissolved 10 g of peptone and 4.0 g of glucose and the resulting solution was adjusted to pH 6.0, after which a drug containing 50 ppb (1% DMSO solution) of each active ingredient was added. Into a 3.5 Ø cm Petri dish was poured 30 ml of the thus obtained Sabouraud medium to prepare an agar plate. Then, the inoculum of the fungus precultured was poured onto the plate in an amount of 0.1 ml per Petri dish. Thus prepared plates were incubated at 28°C for 4-6 days. The test was carried out in duplication for each compound and the results per dish evaluated macroscopically are shown in Table 2.

Evaluation criterion

- 50 + Multiplication of cells was completely inhibited.
- 50 + Multiplication of cells was inhibited.
- 55 - White colonies were formed.
- 55 - The diameter of white colonies increased.

Table 2

Compound No.	Degree of growth	
4	±	+
28	±	+
33	±	±
35	±	+
36	±	+
37	±	±
38	+	+
39	±	±
40	±	±
41	+	++
42	+	±
43	±	+
46	±	±
52	±	±
53	±	+
59	±	±
64	+	+
65	+	+
Control (no compound added)		

Test Example 2

Curing effect test against trichophytosis which was artificially infected in guinea pigs

Hartley strain white male guinea pigs (400 to 600 g) were used as test animals. The hair in three

50 spots on the back of each guinea pig was sheared and then removed in a circle having a diameter of about 3 cm by use of depilatory cream, after which the skin in the depilated spots was lightly rubbed with sandpaper. Thus treated spots were inoculated with 0.1 ml (10^6 spores/spot) of culture of *Trichophyton mentagrophytes* IFO-5466 strain grown on Sabouraud glucose agar media. Each test drug prepared by using polyethylene glycol

300 as base was applied to the inoculation spots once daily for 11 consecutive days in an amount of 0.2 ml per inoculation spot, starting 72 hours after the inoculation. The evaluation was conducted by macroscopical judgement and reverse culture test.

a) Macroscopical judgement:

After the inoculation, the alleviation and development of symptoms in the above-mentioned spots were observed every day for 15 days. The results are shown in Table 3.

- 0: No symptom was observed.
- 1: A few small erythemas were observed.
- 2: Erythemas were insularly scattered or fused into one, and rubefaction was observed around them.

3: Scales were observed, and then formation of thick lesion was observed.

4: Lesion reached to an extreme stage with bleeding.

5 The results obtained are shown in Table 3.

b) Reverse culture test

10 Each guinea pig was sacrificed 15 days after the inoculation, after which the skin in the whole inoculation spots was cut off and the skin cut-off was further cut into small pieces (about 5 mm square). The pieces thus prepared were placed on Sabouraud glucose agar medium containing 20 i.u./ml of penicillin G and 40 µg/ml of streptomycin, and then cultured at 27°C for 2 weeks. The evaluation was carried out by investigating the colonies. The results obtained are shown in Table 4.

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Table 3 (Macroscopic judgement)

Drug tested (Concentration)	Days	4	5	6	7	8	9	10	11	12	13	14	15
Compound 53 (1%)	0.5	1	1	1	1	1	1	1	1	1.5	1.5	2.5	2
Compound 35 (1%)	0.18	1	2	2	2.5	3	3	3.5	3.5	3.5	3.5	3.5	3.5
Compound 38 (1%)	0.18	1	1	1	1	1	1.5	1.5	1.5	1.5	1.5	3	3
Tolnaftate (1%)	0.5	1	1.5	1.5	2	2.5	3	3.5	3.5	3.5	3.5	3.5	4
Compound 38 (0.18%)	0.5	1	1.5	1.5	1	1	1.5	2	2	2	2	2	2
Tolnaftate (1%)	0.18	1	1	1	1.5	2	2.5	3	3	3.5	3.5	3.5	4
Compound 53 (0.18%)	0	1.5	1.5	2	2.5	3	3	3	3	3	3	3	3
Control (No compound treated)	1	2	2	2	3	3.5	3.5	3.5	3.5	4	4	4	4

Table 4 (Results of reverse culture test)

Drug Tested	Concentration	1%	0.1%
Compound 53		0/20	7/20
Compound 35		0/20	6/20
Compound 38		0/20	6/20
Tolnaftate		0/20	16/20
Control (No compound treated)			20/20

Test Example 3

Controlling Effects on Powdery Mildew of Barley

Young barley plants (variety: Kanto No. 6, 2-leaf stage) cultivated in the porcelain pots (12 cm in diameter) were inoculated by conidia of Erysiphe graminis graminis F. sp. hordi, the causal fungus of powdery mildew. The day after inoculation, the plants were treated on the turn table by spraying solutions (200 ppm) of the present compounds by spray gun. The treated plants were kept at 25°C for 6 days in a greenhouse and the development of

lesions was assessed. The controlling effects of the compounds were calculated in comparison with the untreated plots.

The criteria for the controlling effects are:

4; 100 - 95% control

3; 94 - 80% control

2; 79 - 60% control

1; 59 - 0% control

The results are shown in Table 5.

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Table 5

Compound No.	Concen- tration (ppm)	Control- ling effects	Compound No.	Concen- tration (ppm)	Control- ling effects
1	200	3	38	200	4
3	"	4	42	"	4
5	"	4	43	"	4
8	"	4	45	"	4
9	"	4	46	"	4
12	"	3	47	"	4
15	"	4	48	"	4
17	"	4	50	"	4
20	"	4	53	"	4
21	"	4	54	"	4
22	"	4	56	"	4
24	"	4	57	"	4
25	"	4	58	"	4
26	"	4	61	"	4
27	"	2	62	"	4
28	"	4	64	"	4
29	"	4	65	"	4
30	"	2	67	"	4
32	"	2	72	"	4
33	"	4	75	"	4
34	"	4	76	"	4
35	"	4	77	"	4
36	"	4	78	"	4
37	"	4	79	"	4

Test Example 4

Controlling Effects on seed-borne Fusarium

Cucumber seeds infested with Fusarium oxysporum f.sp. cucumerinum were soaked in solutions (200 ppm) of the present compounds for 20

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hr. The treated seeds were then placed on the selective medium for Fusarium spp. (Komada medium). After a week-incubation at 25°C, the mycelial growth around the treated seeds was assessed and the controlling effects were calculated in comparison with untreated plots. The criteria for controlling effects are as same as in Test Example 3. The results are shown in Table 6.

Table 6

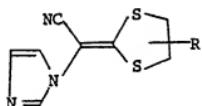
Compound No.	Concen- tration (ppm)	Control- ling effects	Compound No.	Concen- tration (ppm)	Control- ling effects
3	200	4	52	200	4
4	"	4	53	"	4
6	"	4	54	"	4
7	"	4	55	"	4
10	"	4	56	"	4
11	"	4	57	"	4
13	"	4	58	"	3
16	"	4	60	"	4
18	"	4	61	"	4
20	"	4	63	"	4
21	"	4	64	"	4
24	"	3	65	"	4
26	"	2	66	"	4
29	"	4	67	"	2
33	"	4	68	"	4
35	"	4	69	"	4
37	"	4	70	"	4
39	"	4	71	"	4
40	"	4	73	"	4
44	"	4	75	"	4
47	"	4	76	"	4
49	"	4	77	"	4
50	"	4	78	"	4
51	"	4	79	"	4

The recipes shown below as Formulation Examples 1 to 3 are for medical purposes. For these purposes, the various adjuvants and constituents to

be used should be of pharmaceutically acceptable grade. In Formulation Examples 1 to 3, all parts are

by weight.	Compound 12 50 parts
Formulation Example 1	Mixture of diatomaceous earth and clay 45 parts
Compound 35 1 part	5 Polyoxyethylene nonylphenyl ether 5 parts These ingredients are homogeneously mixed and then pulverized to prepare a wettable powder.
Polyethylene glycol 300 95 parts These ingredients are mixed to prepare an emetic solution.	70 Formulation Example 5
Formulation Example 2	Emulsifiable concentrate
Compound 53 2 parts	75 Compound 23 20 parts
Polyethylene glycol 400 40 parts	Tetrahydrofuran 20 parts
Polyethylene glycol 1500 58 parts These ingredients are mixed with heating to obtain a solution, which is then cooled to prepare an ointment.	Xylene 45 parts 20 Mixture of polyoxyethylene nonylphenyl ether and alkyl benzenesulfonate 15 parts These ingredients are homogeneously mixed to prepare an emulsifiable concentrate.
Formulation Example 3	25 Formulation Example 6
Compound 38 2 parts	Dust
1,2-Propanediol 5 parts	30 Compound 29 4 parts
Glycerol stearate 5 parts	Mixture of diatomaceous earth, clay and talc 95 parts
Spermaceti 5 parts	35 Calcium stearate 1 part These ingredients are homogeneously mixed and then pulverized to prepare a dust.
Isopropyl myristate 10 parts	40 Formulation Example 7
Polyisobutene 60 4 parts A mixture of them is heated and then cooled, after which 69 parts of water is added with stirring to prepare cream.	Granules
In addition to the pharmaceutical formulations described above, preparation in pharmaceutically usable forms such as injections, tablets and the like is possible.	45 Compound 46 3 parts
The recipes shown in Formulation Examples 4 to 7 are for agricultural chemicals. In these examples, all parts are by weight as in Formulation Examples 1 to 3.	Mixture of bentonite and clay 92 parts
Formulation Example 4	50 Calcium lignin sulfonate 5 parts These ingredients are homogeneously mixed and then pulverized, after which an adequate amount of water is added, and the resulting mixture is sufficiently kneaded and then granulated to prepare granules.
Wettable powder	55 Claims

1. A ketene S,S-acetal derivative represented by the general formula (I):

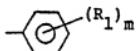


(I)

wherein R represents a hydrogen atom; an alkyl group having 1 to 8 carbon atoms; a cycloalkyl group having 3 to 6 carbon atoms; a methylene group; a lower alkynyl group; a lower alkyl group

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substituted by a halogen atom, a cyano group, a lower alkoxy group, a lower alkylthio group, a carbamoyl group, an acyl group, or an alkenyloxy groups; a phenyl group represented by



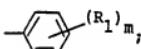
(in which R₁ represents a hydrogen atom, a halogen atom, a straight or branched chain lower alkyl group, a lower alkoxy group which may be substituted by one or more halogen atoms, a phenoxy group or a methylenedioxy group, and m represents an integer of 1 to 3); a benzyl group; a methylenedioxybenzyl group; a phenoxyalkyl

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group; a phenoxyalkyl group substituted by a halogen atom; a naphthyl group; or a substituted or unsubstituted pyridyl group.

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2. A ketene S,S-acetal derivative according to Claim 1, wherein R represents a lower alkyl group, a lower alkyl group substituted by a lower alkoxy or alkylthio group, or

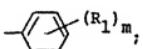


in which R₁ represents a halogen atom, or a straight or branched chain lower alkyl group, and m represents an integer of 1 to 3.

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3. A ketene S,S-acetal derivative according to Claim 1, wherein R represents

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in which R₁ represents a halogen atom or methyl group, and m represents an integer of 1 to 3.

4. A ketene S,S-acetal derivative according to Claim 1, which is 2-(1-imidazolyl)-2-[4-(2,4-dichlorophenyl)-1,3-dithiolan-2-ylidene]acetonitrile (E isomer).

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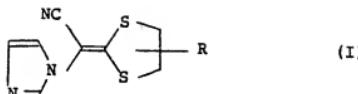
5. A ketene S,S-acetal derivative according to Claim 1, which is 2-(1-imidazolyl)-2-[4-(2-chlorophenyl)-1,3-dithiolan-2-ylidene]acetonitrile (E isomer).

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6. A ketene S,S-acetal derivative according to Claim 1, which is 2-(1-imidazolyl)-2-[4-(2-bromo-phenyl)-1,3-dithiolan-2-ylidene]acetonitrile (E isomer).

7. A ketene S,S-acetal derivative according to Claim 1, which is 2-(1-imidazolyl)-2-[4-(4-chlorophenyl)-1,3-dithiolan-2-ylidene]acetonitrile (E isomer).

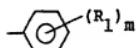
8. A process for producing a ketene S,S-acetal derivative represented by the general formula (I):



wherein R, represents a hydrogen atom; an alkyl group having 1 to 8 carbon atoms; a cycloalkyl group having 3 to 6 carbon atoms; a methylene group; a lower alkenyl groups; a lower alkyl group

substituted by a halogen atom, a cyano group, a lower alkoxyl group, a lower alkylthio group, a carbamoyl group, an acyl group, or an alkenyloxy group; a phenyl group represented by

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(in which R, represents a hydrogen atom, a halogen atom, a straight or branched chain lower alkyl group, a lower alkoxy group which may be substituted by one or more halogen atoms, a phenoxy group or a methylenedioxy group, and m represents an integer of 1 to 3); a benzyl group; a

methylenedioxybenzyl group; a phenoxyalkyl group; a phenoxyalkyl group substituted by a halogen atom; a naphthyl group; or a substituted or unsubstituted pyridyl group, which comprises reacting 1-cyanomethylimidazole represented by the structural formula (II):

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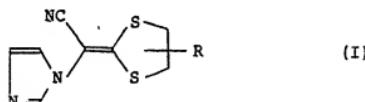
with carbon disulfide in the presence of a base, and then reacting the reaction product with a compound represented by the general formula (III):



wherein R is as defined above, and X represents a halogen atom, a mesyloxy or a tosyloxy group.

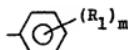
30 9. A method for curing mycosis which comprises administering a pharmaceutical preparation containing a pharmaceutically effective amount of a ketene S,S-acetal derivative represented by the general formula (I):

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wherein R represents a hydrogen atom; an alkyl group having 1 to 8 carbon atoms; a cycloalkyl group having 3 to 6 carbon atoms; a methylene group; a lower alkenyl group; a lower alkyl group

50 substituted by a halogen atom, a cyano group, a lower alkoxyl group, a lower alkylthio group, a carbamoyl group, an acyl group, or an alkenyloxy group; a phenyl group represented by



(in which R₁ represents a hydrogen atom, a halogen atom, a straight or branched chain lower alkyl group, a lower alkoxy group which may be substituted by one or more halogen atoms, a phenoxy group or a methylenedioxy group, and m represents an integer of 1 to 3); a benzyl group; a methylenedioxybenzyl group; a phenoxyalkyl group; a phenoxyalkyl group substituted by a halogen atom; a naphthyl group; or a substituted or unsubstituted pyridyl group.

10. A method for curing mycosis according to Claim 9, wherein the ketene S,S-acetal derivative is 2-(1-imidazolyl)-2-[4-(2,4-dichlorophenyl)-1,3-dithiolan-2-ylidene]acetonitrile (E isomer).

11. A method for curing mycosis according to Claim 9, wherein the ketene S,S-acetal derivative is 2-(1-imidazolyl)-2-[4-(2-chlorophenyl)-1,3-dithiolan-2-ylidene]acetonitrile (E isomer).

12. A method for curing mycosis according to Claim 9, wherein the ketene S,S-acetal derivative is 2-(1-imidazolyl)-2-[4-(2-bromophenyl)-1,3-dithiolan-2-ylidene]acetonitrile (E isomer).

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PARTIAL EUROPEAN SEARCH REPORT

which under Rule 45 of the European Patent Convention
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Application number

EP 85 11 2797.7

DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int. Cl. *)
A	<u>EP - A - 0 080 103 (BAYER AG)</u> * claims 1, 3 *	1,8	C 07 D 409/06 C 07 D 409/14 A 01 N 43/50
A	<u>DD - A - 202 713 (M. AUGUSTIN et al.)</u> * example 2 *	1,8	
A	<u>EP - A - 0 061 794 (JANSSEN PHARMACEUTICAL N.V.)</u> * page 1, line 18 - page 2, line 12 *	1	
A	<u>DE - A - 2 134 499 (BAYER AG)</u> * claim 1 *	1	

TECHNICAL FIELDS
SEARCHED (Int. Cl. *)

C 07 D 409/00
C 07 D 521/00
A 01 N 43/00

INCOMPLETE SEARCH

The Search Division considers that the present European patent application does not comply with the provisions of the European Patent Convention to such an extent that it is not possible to carry out a meaningful search into the state of the art on the basis of some of the claims.

1-5

Claims searched completely:

9-12

Claims searched incompletely:

9-12

Claims not searched:

9-12

Reason for the limitation of the search:

9-12 (Method for treatment of
the human or animal body
by therapy; Art. 52.4)

Place of search	Date of completion of the search	Examiner
Berlin	23-05-1986	VAN AMSTERDAM
CATEGORY OF CITED DOCUMENTS		
X : particularly relevant if taken alone	T : theory or principle underlying the invention	
Y : particularly relevant if combined with another document of the same category	E : earlier patent document, but published on, or after the filing date	
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